

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
15 March 2001 (15.03.2001)

PCT

(10) International Publication Number
WO 01/18822 A1

(51) International Patent Classification: G21C 21/02, 3/62

(21) International Application Number: PCT/EP00/08057

(22) International Filing Date: 17 August 2000 (17.08.2000)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
99116886.5 6 September 1999 (06.09.1999) EP

(71) Applicant (for all designated States except US): EURO-
PEAN COMMUNITY (EC) [—/LU]; Rue Alcide de
Gasperi, L-2920 Luxembourg (LU).

(72) Inventors; and

(75) Inventors/Applicants (for US only): HAAS, Didier

[BE/DE]; Eisbergweg 12, D-76356 Weingarten (DE).
FUCHS, Claude [FR/FR]; 4, place de la Chapelle,
F-67360 Niederlauterbach (FR). FOURCAUDOT,
Serge [FR/DE]; Karlsstr. 99, D-76137 Karlsruhe (DE).
CHAROLLAIS, François [FR/DE]; Bahnhofstr. 16b,
D-76351 Linkenheim (DE). SOMERS, Joseph [IE/DE];
Reinhold-Frank-Str. 4, D-76133 Karlsruhe (DE).

(74) Agent: WEINMILLER, Jürgen; Spott & Weinmiller,
Lennéstr. 9, D-82340 Feldafing (DE).

(81) Designated States (national): CA, JP, NO, US.

Published:

— With international search report.

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: METHOD FOR PRODUCING NUCLEAR FUEL PELLETS OF THE MOX TYPE

(57) Abstract: A method for producing nuclear fuel pellets of the MOX (mixed plutonium and uranium oxide) type, comprising the steps of preparing an U-pu oxide blend powder having a Pu content in excess of the finally desired value, preparing a uranium oxide powder, mixing adequate quantities of both powders in order to achieve the desired plutonium content, compacting and sintering the mixture for obtaining said pellets, wherein the step of preparing the uranium oxide powder involves the following sequence of substeps: a) preparation of an aqueous solution of uranyl nitrate to which between 0.5 and 2 wt% of organic thickeners are added such that the viscosity of the solution is adjusted to values between 20 and 100 centipoise, b) dispersion of the solution into droplets, c) introducing said droplets into a hydroxide bath, d) washing the resulting beads, e) drying the beads by azeotropic distillation using an immiscible organic solvent, f) thermal treatment of the beads in an oxidising atmosphere, g) thermal treatment in a reducing atmosphere.

WO 01/18822 A1